

Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) **EP 1 166 854 A1**

(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:
02.01.2002 Bulletin 2002/01

(51) Int Cl.7: **B01D 53/94**

(21) Application number: 01114550.5

(22) Date of filing: 18.06.2001

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR**
Designated Extension States:
AL LT LV MK RO SI

(72) Inventors:
• Matsumoto, Takeshi
Himeji-shi, Hyogo 670-0971 (JP)
• Horiuchi, Makoto
Himeji-shi, Hyogo 671-2221 (JP)
• Yoshikawa, Tatsuya
Himeji-shi, Hyogo 670-0052 (JP)

(30) Priority: 20.06.2000 JP 2000185257

(71) Applicants:
• ICT Co., Ltd.
Osaka-shi, Osaka-fu (JP)
• International Catalyst Technology, Inc.
Ridgefield Park, N.Y. 07660 (US)

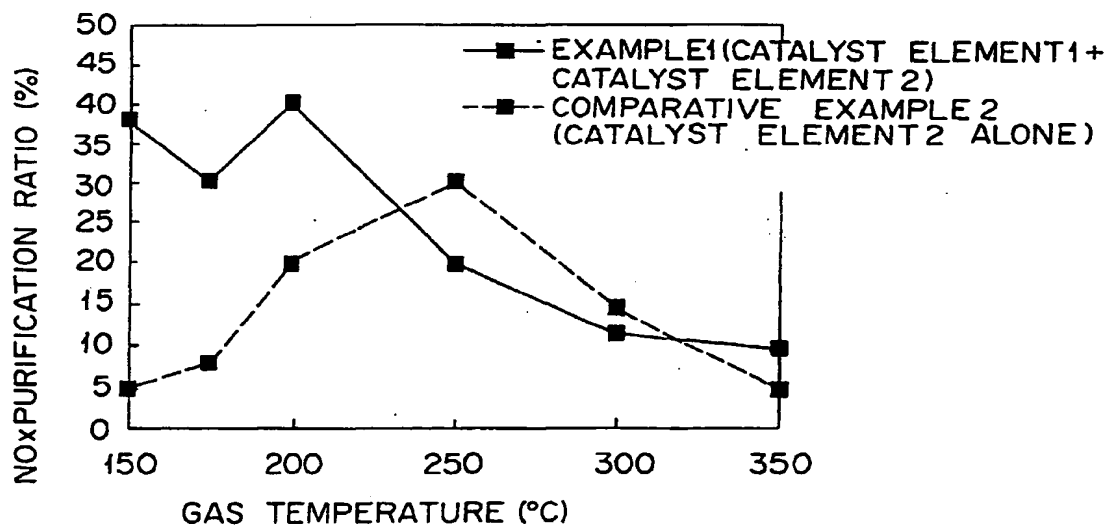
(74) Representative:
Luderschmidt, Schüler & Partner GbR
Patentanwälte, John-F.-Kennedy-Strasse 4
65189 Wiesbaden (DE)

(54) **Exhaust gas purifying catalyst and method for purifying exhaust gas**

(57) An exhaust gas purifying catalyst including a catalyst component containing a refractory inorganic oxide carrying a platinum family metal, a nitrogen oxide

adsorbent, and a hydrocarbon adsorbent and a catalyst component for the purification of nitrogen oxide and a method for the purification of an exhaust gas by the use of the catalyst.

FIG.2



EP 1 166 854 A1

Description

BACKGROUND OF THE INVENTION

Field of the Invention:

[0001] This invention relates to an exhaust gas purifying catalyst and a method for the purification of exhaust gas. More particularly, it relates to a method for catalytically reducing harmful substances, particularly nitrogen oxides (NO_x), contained in the exhaust gas with a hydrocarbon as a reducing agent, an exhaust gas purifying catalyst capable of adsorbing simultaneously the hydrocarbon and NO_x in the exhaust gas thereby decomposing and reducing NO_x in a wide temperature range without introducing any hydrocarbon by way of replenishment from the external source, and a method for the purification of the exhaust gas by the use of the catalyst.

Description of Related Art:

[0002] The NO_x in the atmospheric air causes photochemical smog and acid precipitation. Thus, the emission of NO_x from mobile generative sources such as automobiles that are furnished with internal combustion engines such as gasoline and diesel engines, which form one of the generative sources for NO_x , has become one of the serious social issues. Official studies are now under way with a view to rigidifying legal regulations on the tolerance of NO_x emission in the future. In the light of the demand of such social cause, efforts are being promoted with a view to developing an exhaust gas purifying catalyst.

[0003] As the catalyst capable of purifying the exhaust gas of a gasoline engine, the so-called three-way catalyst, i. e. a catalyst capable simultaneously reducing NO_x , unburnt hydrocarbon, and carbon monoxide, has been heretofore used. The three-way catalyst, when the ordinary gasoline engine is used and the exhaust gas therefrom contains substantially no oxygen, is capable of efficiently reducing NO_x with the unburnt hydrocarbon and the carbon monoxide in the exhaust gas.

[0004] The exhaust gas from the diesel engine contains oxygen in great excess on account of the special quality of the engine, and theoretically has small contents of hydrocarbon and carbon monoxide as a reducing agent for NO_x . When the ordinary three-way catalyst is used, it is nearly incapable of reducing NO_x for the treatment of exhaust gas from the diesel engine.

[0005] In recent years, gasoline engines of type of a lean burn and an in-cylinder fuel injection system have been developed. Since these engines operate by lean burn of the fuel, their exhaust gases contain oxygen in high concentrations. When the ordinary three-way catalyst is used, it purifies NO_x with difficulty for the treatment of such an exhaust gas of engines.

[0006] JP-A-63-100919 proposes a catalyst having copper deposited on such a porous carrier as zeolite and JP-A-5-137963 proposes a method using platinum as a main catalyst as respects the exhaust gas purifying catalyst effective in the removal NO_x in the exhaust gas containing oxygen copiously such as the exhaust gas of a diesel engine or a lean burn gasoline engine.

[0007] These methods heretofore proposed have, as conditions for attaining efficient decomposition of NO_x , the point that at least the temperature is limited in a comparatively narrow range and the point that the amount of the hydrocarbon as a reducing agent is comparatively large. These conditions are not easily applied to the actual exhaust gas that varies with the kind of engine and the conditions of travel.

[0008] Not only the copper and platinum type catalysts mentioned above but also the catalyst in general is incapable of purifying NO_x unless the catalyst bed is heated to a certain degree by the exhaust gas from the engine. In the diesel engine whose exhaust gas has a comparatively low temperature, it is extremely difficult for the conventional technique to purify NO_x that is generated from immediately after the start of the engine through the duration of idling and low-speed travel.

[0009] From this view, a NO_x adsorbent that is not appreciably affected by the exhaust gas temperature and the gas composition as compared with NO_x purifying catalyst has been proposed. Specifically, JP-A-9-57093 discloses an adsorbent having nickel oxide and copper oxide as main components and aimed at the NO_x contained in the atmospheric air or in various kinds of roadside exhaust gas, and JP-A-7-47227 discloses an adsorbent having iron oxide hydrate as a main component and aimed particularly at NO_x contained in the exhaust gas in an automobile tunnel or an indoor parking lot. The NO_x adsorbents thus proposed, however, have limited capacities for adsorption, are incapable of being continuously used and needed regeneration as an indispensable condition. Further, these NO_x adsorbents are generally used under conditions such as a fixed temperature and a gas composition variable only sparingly. Even when NO_x adsorbent is used on the exhaust gas from a diesel engine, since the exhaust gas is always in a lean atmosphere, the desorption of NO_x by the reducing gas and the regeneration of the reducing agent by purification cannot be hoped for. Further, since NO_x adsorbent has the behavior of adsorption and desorption of NO_x largely de-

pending on temperature, it entails the problem that NO_x adsorbed in a low temperature range, on exposure to a high temperature, is inevitably desorbed without being purified.

SUMMARY OF THE INVENTION

[0010] This invention has been produced in view of the conventional problems mentioned above.

[0011] An object of the present invention is to provide an exhaust gas purifying catalyst capable of efficiently removing NO_x in the exhaust gas of diesel engine that usually retains a lean atmosphere, has a small content of hydrocarbon as a reducing gas, and is under a low temperature.

[0012] Another object of the present invention is to provide a method for purifying the exhaust gas by the use of this catalyst.

[0013] We have pursued various studies one after another in search of an exhaust gas purifying catalyst with a view to accomplishing the object described above, and have found that the problems mentioned above can be solved by combining a refractory inorganic oxide carrying thereon a platinum family metal, a catalyst containing a NO_x adsorbent and a hydrocarbon (HC) adsorbent, and a NO_x purifying catalyst. This invention has been perfected as a result.

[0014] The object of this invention are accomplished by an exhaust gas purifying catalyst that is characterized by combining a first catalyst component containing a refractory inorganic oxide carrying thereon a platinum family metal, a NO_x adsorbent, and a HC adsorbent and a second catalyst component of a NO_x purifying catalyst.

[0015] Another object of this invention is accomplished by a method for the purification of an exhaust gas, characterized by forwarding an exhaust gas having a hydrocarbon and a NO_x at a molar ratio (hydrocarbon: nitrogen oxide) in the range of 0.1 to 2: 1 into contact with the catalyst above.

[0016] According to the invention, in the method for catalytic reduction of the harmful substances, particularly NO_x, contained in the exhaust gas with a hydrocarbon as a reducing agent, it is capable of adsorbing simultaneously HC and NO_x in the exhaust gas thereby decomposing or reducing NO_x in a wide temperature range without introducing a large amount of a hydrocarbon by way of replenishment from the external source. It is particularly effective in reducing HC and NO_x in low catalyst temperatures as when the engine is started.

[0017] The above and other objects, features and advantages of the present invention will become clear from the following description of the preferred embodiments.

BRIEF DESCRIPTION OF DRAWINGS

[0018] The accompanying drawing incorporated in and forming a part of the specification, illustrates several aspects of the present invention, and together with the description serve to explain the principles of the invention. In the drawings:

Fig.1 is a graph showing the special quality of adsorption and emission of NO_x and HC demonstrated by the catalyst in Comparative Example 1;

Fig. 2 is a graph showing purification ratios of NO_x demonstrated by the catalysts of Example 1 and Comparative Example 2; and

Fig. 3 is a graph showing purification ratios of NO_x demonstrated by the catalysts of Example 1 and Comparative Examples 3 and 4.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0019] This invention uses a nitrogen oxide (NO_x) adsorbent and a hydrocarbon (HC) adsorbent, and is capable of effectively reducing the harmful substances in the exhaust gas by adsorbing NO_x and HC when the catalyst temperature is low as during the course extending from immediately after the start of engines through the phase of idling. Further, when the temperature is high, since the adsorbed NO_x and HC are desorbed, NO_x can be effectively reduced with the NO_x purifying catalyst utilizing the desorbed HC as a reducing agent.

[0020] Now, this invention will be described more specifically below.

[0021] The NO_x adsorbent is a main component forming a first catalyst component, and commendable to use nickel oxide or α -hydrated iron oxide. Though nickel oxide occurs in various states, it is chiefly sole Ni₂O₃ or a mixture of NiO and Ni₂O₃. For this invention, the presence of Ni₂O₃ is indispensable from the viewpoint of the quality of NO_x adsorption. If nickel oxide formed solely of NiO is used, the produced catalyst will not acquire a satisfactory ability to adsorb NO_x. Then, the decomposition temperature of Ni₂O₃ is 350°C. When nickel oxide is adopted as a NO_x adsorbent, it is commendable to use this catalyst at a temperature below the decomposition temperature. The hydrated iron oxide is known in three kinds, α , β , and γ , which differ in crystal structure. The variations that have respective OH⁻ ions and O²⁻ ions arranged characteristically on the crystal faces thereof may be cited. The hydrated iron oxide is thermally instable and,

on being exposed to an elevated temperature, becomes stable as iron oxide (Fe_2O_3) with release of water. When the hydrated iron oxide is adopted as a NO_x adsorbent, it is commendable to use this adsorbent at a temperature of not higher than 500°C . In this invention, the α -hydrated iron oxide is used favorably.

[0022] The amount of the NO_x adsorbent to be used may comprise from 10 to 100 g, as reduced to oxide, per liter of the catalyst. If this amount falls short of 10 g, the shortage will be at a disadvantage in exhibiting an insufficient ability to adsorb NO_x and, as a result, narrowing the temperature range which allows NO_x purification. Conversely, if the amount exceeds 100 g, the excess will proportionately add to the amount of NO_x to be adsorbed but will nevertheless be at a disadvantage in unduly increasing the amount of NO_x to be desorbed when the temperature of the exhaust gas is elevated. The NO_x purifying catalyst, therefore, does not purify them fully but rather suffers the ability to purify NO_x to decline. When this material is slurried, the produced slurry entails degradation of its quality.

[0023] The HC adsorbent that forms the first catalyst component is preferred to be zeolites. It is commendable for the zeolites to have a high ability to adsorb hydrocarbon and excel in heat resistance. Examples of the zeolites may include Pentasil type zeolite, Y type zeolite, mordenite, and ferrierite. The amount of HC adsorbent to be used may comprise from 10 to 100 g per liter of the catalyst. If the amount falls short of 10 g, the shortage will be at a disadvantage in preventing the adsorbent from manifesting a satisfactory ability to adsorb the hydrocarbon. Conversely, if the amount exceeds 100 g, the excess will proportionately increase the amount of the hydrocarbon to be adsorbed but will be nevertheless at a disadvantage in preventing the NO_x purifying catalyst from acquiring a proportionate exaltation of the ability thereof. When the material is slurried, the produced slurry will degrade its quality.

[0024] The platinum family metal contained in the first catalyst component is at least one member selected from the group consisting of platinum, palladium, rhodium and mixtures thereof. Platinum and palladium prove particularly favorable from the viewpoint of the treatment of NO_x . Such a metal is generally used as deposited on the powder of a refractory inorganic oxide such as alumina, silica, silica-alumina, zirconia, titania, and zeolite which have a large specific surface area. Titania proves particularly advantageous from the viewpoint of heat resistance. The deposition of Noble metal on the powder is accomplished by adopting any of suitable known methods such as ion exchange or impregnation method. The amount of the platinum family metal to be used is not restricted but may be fixed depending on the kind of metal to be used and the reaction conditions under which the catalyst is placed. It may comprise from 0.001 to 1 g, preferably from 0.01 to 0.5 g, as reduced to metal, per liter of the catalyst. If the amount falls short of 0.001 g, the shortage will be at a disadvantage in preventing the catalyst to acquire an unsatisfactory ability to oxidize and entailing the problem of coking. Conversely, if the amount exceeds 1 g, the excess will promote the combustion of the hydrocarbon and inevitably degrade the NO_x purification ratio because of the insufficient supply of the hydrocarbon necessary for the NO_x purification.

[0025] Then, the amount of the refractory inorganic oxide to be used may comprise from 2 to 50 g, preferably from 5 to 20 g per 1 g of the platinum family metal. If this amount falls short of 2 g, the shortage will be at a disadvantage in inducing insufficiency of dispersibility during colorization. Conversely, if this amount exceeds 50 g, the excess will be at a disadvantage in aggravating the property.

[0026] A second catalyst component for the NO_x purification is not restricted so long as it begins to purify NO_x from a low temperature upward. The catalyst that is formed by depositing a Pt material on a porous inorganic oxide such as zeolites proves advantageous. The zeolite is preferred to manifest high activity after ion-exchanging Pt and excels in heat resistance. Examples of the zeolites may include Pentasil type zeolite, Y type zeolite, mordenite, and ferrierite. Further, when a porous inorganic oxide such as zeolites is deposited on a refractory three-dimensional structure, it is commendable to use an inorganic binder such as silica sol to create firm adhesion in the interface in an appropriate amount, preferably in the range of 0.01 to 0.5 g as a solid component per g of the zeolite.

[0027] The amount of the first catalyst component may comprise from 10 to 250 g per liter of the catalyst.

[0028] The amount of the second catalyst component may comprise from 10 to 300 g, preferably from 50 to 150 g, per liter of the catalyst. The amount of Pt contained therein is generally in the range of 0.01 to 2 g, preferably in the range of 0.1 to 1 g, per liter of the catalyst. If this amount falls short of 0.01 g, the shortage will be at a disadvantage in causing only an insufficient reaction between NO_x and the hydrocarbon and consequently attaining no sufficient ratio of NO_x purification. Conversely, if this amount exceeds 2 g, the excess will be at a disadvantage in promoting the combustion of the hydrocarbon and inevitably degrading the NO_x purification ratio.

[0029] In this invention, these first and second catalyst components may be used as deposited on a refractory three-dimensional structure.

[0030] The refractory three-dimensional structure may include an integrally molded honeycomb structure such as a honeycomb carrier, a monolithic honeycomb carrier, a metal honeycomb carrier, and a plug honeycomb carrier.

[0031] The monolithic carrier may be what is generally called a honeycomb carrier. Particularly, honeycomb carriers using cordierite, mullite, α -alumina, zirconia, titania, titanium phosphate, aluminum titanate, betalite, spodumene, aluminosilicate, and magnesium silicate as raw materials prove favorable. Those made of cordierite proves especially advantageous. Besides, the refractory three-dimensional structures formed of antioxidant refractory metals such as stainless steel and Fe-Cr-Al alloy are also usable.

[0032] The monolithic carrier may be produced by methods of extrusion molding or tightly rolling a sheet-like element. The gas-passing openings (cells) in the monolithic carrier may be in a hexagonal, rectangular, triangular, or corrugated shape. For satisfactory service of the monolithic carrier, the cell density (the number of cells per unit cross-sectional area, 6.45 cm² (1 square inch)) may be in the range of 100 - 600 cells, preferably in the range of 200 - 500 cells.

[0033] The term "catalyst" as used in this invention embraces one or a combination of a plurality of (catalyst elements) having the first catalyst component and/or the second catalyst component deposited on the structure. The catalyst element may include one formed by depositing both first and second catalyst components on one refractory three-dimensional structure or a combination thereof or one formed by separately depositing both first and second catalyst components on each of the structures and then combining them.

[0034] The method for producing the catalyst element is not restricted but may cite any of the known methods. Example of the method for depositing the first and second catalyst components on the structure may include the following method.

[0035] As respects the first catalyst component, a powder containing platinum is obtained by immersing the powder of an oxide of titanium in the aqueous solution of a salt of platinum, for example, drying the immersed powder at a temperature of 80°C to 250°C, and calcining the dried powder at a temperature of 300°C to 850°C for a period of 0.5 to 5 hours.

[0036] The produced powder, a NO_x adsorbent such as a nickel oxide powder, and a HC adsorbent such as zeolite are slurried by wet pulverizing with a wet pulverizer such as a ball mill. An open flow type honeycomb carrier made of cordierite, as a refractory three-dimensional structure, is immersed in the produced slurry. Subsequently, the honeycomb carrier nowwet with the slurry is drained to expel excess slurry and then dried at a temperature of 80°C to 250°C for a period of 0.5 to 5 hours.

[0037] As respects the second catalyst component, a powder is obtained by immersing a porous carrier such as zeolite in the aqueous solution of a salt of platinum, for example, drying the wet carrier at a temperature of 80°C to 250°C, preferably of 100°C to 150°C, and calcining the dried carrier at a temperature of 300°C to 850°C, preferably of 400°C to 700°C, for a period of 0.5 to 5 hours, preferably of 1 to 2 hours. When all the oxide powder excepting noble metals is used, it may be subjected, as it is, to the subsequent wet pulverizing step without drying and calcining.

[0038] Next, the powder of platinum and zeolite, a binder such as silica sol, and water are wet pulverized with a wet pulverizer such as a ball mill till they form slurry. An open flow type honeycomb made of cordierite, as the refractory three-dimensional structure, is immersed in the produced slurry. The wet honeycomb is drained to expel excess slurry, then dried at a temperature of 80°C to 250°C, preferably of 100°C to 150°C, and, when necessary, calcined at a temperature of 300°C to 800°C, preferably of 400°C to 700°C, for a period of 0.5 to 3 hours, preferably of 1 to 2 hours.

[0039] It is, of course, permissible to prepare slurry containing these first and second catalyst components and then immerse a refractory three-dimensional structure in the slurry obtained.

[0040] The catalyst components obtained as described above are not restricted in their disposition or composition. They may be disposed by methods heretofore known to the art, but may include the following methods. (1) The first catalyst component is disposed on the upstream side and the second catalyst component is disposed on the downstream side relative to the flow direction of the exhaust gas. (2) The first and second catalyst components are uniformly mixed and then the produced mixture is applied to the structure. (3) The first catalyst component is made to form the inner layer of a catalyst bed and the second catalyst component to form the outer layer of the catalyst bed. In the above method (1), when the first and second catalyst components are deposited on one refractory three-dimensional structure, namely when one catalyst element is used, the method of deposition may adopted which causes the concentration of the first catalyst component to be gradually lowered from the upstream side through the downstream side relative to the exhaust gas flow and conversely the concentration of the second catalyst component to be gradually heightened. It is, of course, permissible to deposit only the first catalyst component on the upstream side and only the second catalyst component on the downstream side.

[0041] When a plurality of refractory three-dimensional structures having both the first and second catalyst components simultaneously or separately deposited thereon are used in combination, namely when a plurality of catalyst elements are used, the method may be adopted which uses the catalyst elements having the first and second catalyst components deposited respectively in varying amounts and disposes the catalyst elements having the first catalyst components deposited in a large amount on the upstream side and the catalyst elements having the second catalyst components deposited in a larger amount on the downstream side respectively relative to the flow direction of the exhaust gas. It is, of course, permissible to adopt the method of disposing the catalyst elements having only the first catalyst component deposited thereon on the upstream side and the catalyst elements having only the second catalyst component deposited thereon on the downstream side, respectively.

[0042] The exhaust gas is preferred to be such that the molar ratio of the hydrocarbon to the nitrogen oxide (hydrocarbon: nitrogen oxide) therein is in the range of 0.1 to 2: 1. An exhaust gas from diesel engines is preferred. Further, during the purification of exhaust gases, the inlet temperature of the exhaust gas purifying catalyst is preferably adjusted in the range of 100°C to 500°C.

EXAMPLES

[0043] Now, this invention will be described more specifically below with examples and comparative examples. It should be noted, however, that this invention is not limited by these examples.

EXAMPLE 1

[0044] A titania powder (available from Ishihara Sangyo Kaisha, Ltd. in Japan) was immersed with an aqueous dinitrodiamineplatinum solution, dried at 120°C, and then calcined at 500°C for one hour to obtain a Pt-deposited titania powder (Powder 1). The Pt concentration of this powder was 10 wt. % based on the weight of titania.

[0045] A quantity 0.22 g of the resultant Powder 1, 1000 g of a nickel oxide powder (reagent grade, available from Wako Pure Chemical Industries, Ltd. in Japan), 1080 g of a ZSM-5 type zeolite (available from zeolyst International), and 2000 g of water were ball milled. In the produced slurry, an open flow type honeycomb carrier formed of cordierite (1.0 liter, 400 cells, available from NGK Insulators Ltd. in Japan) as a refractory carrier was immersed. Subsequently, the honeycomb carrier wet with the slurry was drained to expel excess slurry and then dried at 150°C for two hours. The resulting catalyst element (Catalyst Element 1) contained 0.01 g of Pt, 0.1 g of TiO₂, 50 g of nickel oxide, and 50 g of ZSM-5, per liter of carrier.

[0046] Separately, 1 kg of an aqueous platinum ammine hydroxide solution (Pt: 1.2 g/liter) was mixed with 1200 g of a zeolite powder, stirred, then filtered and the resultant dried at 80°C for 12 hours, and subsequently calcined at 500°C for one hour to afford a Pt ion-exchange zeolite powder (Powder 2). The Pt concentration of this powder was 0.1%.

[0047] A slurry was formed by wet pulverizing 1092 g of Powder 2, 544 g of silica sol (solid content 20%) (available from Nissan Chemicals Industries Ltd. in Japan), and 1032 g of water. A honeycomb carrier (supra) was immersed in the produced slurry. The honeycomb carrier wet with the slurry was drained to expel excess slurry and then dried at 150°C for two hours. The resulting catalyst element (Catalyst Element 2) contained 100.1 g of Pt-ZSM-5 and 10 g of SiO₂, per liter of carrier.

[0048] Disposing Catalyst Element 1 on the former stage (upstream side) and Catalyst Element 2 on the latter stage (downstream side) formed a catalyst.

EXAMPLE 2

[0049] A quantity 0.22 g of Powder 1, 1000 g of a nickel oxide powder (supra), 1080 g of a ZSM-5 type zeolite (supra), and 2000 g of water were ball milled.

[0050] Separately, 1092 g of Powder 2, 544 g of silica gel (supra), and 1032 g of water were ball milled.

[0051] First, a honeycomb carrier (supra) was immersed to one half in the length direction in the slurry made of Powder 1, nickel oxide, and zeolite. Subsequently the honeycomb carrier wet with the slurry was drained to expel excess slurry and then dried at 150°C for two hours.

[0052] Next, in the slurry formed of Powder 2, the produced carrier was immersed on the opposite side, namely the part carrying no catalyst component. The carrier wet with the slurry was drained to expel excess slurry and dried at 150°C for two hours. The resulting catalyst element (Catalyst Element 3) contained 0.01 g of Pt, 0.1 g of TiO₂, 50 g of nickel oxide, 50 g of ZSM-5, 100.1 g of Pt-ZSM-5, and 10 g of SiO₂, per liter of carrier.

[0053] Catalyst Element 3 alone was used as a catalyst and the part formed of Powder 1, nickel oxide, and zeolite was disposed on the upstream side, and the part formed of Powder 2 disposed on the downstream side.

EXAMPLE 3

[0054] A quantity 1.1 g of Powder 1, 500 g of a nickel oxide powder (supra), 540 g of a ZSM-5 type zeolite, 2000 g of water, and 1001 g of Powder 2 were ball milled. A honeycomb carrier (supra) was immersed in the resultant slurry. The honeycomb carrier now wet with the slurry was drained to expel excess slurry and then dried 150°C for two hours. The resulting catalyst element (Catalyst Element 4) contained 0.01 g of Pt, 0.1 g of TiO₂, 50 g of nickel oxide, 50 g of ZSM-5, and 100.1 g of Pt-ZSM, per liter of carrier.

[0055] Catalyst Element 4 alone was used as a catalyst.

EXAMPLE 4

[0056] A quantity 1092 g of Powder 2, 544 g of silica sol (supra), and 1032 g of water were ball milled. Catalyst Element 1 was immersed in the resultant slurry. Subsequently, Catalyst Element 1 now wet with the slurry was drained to expel excess slurry and then dried 150°C for two hours. The resulting catalyst element (Catalyst Element 5) contained

EP 1 166 854 A1

0.01 g of Pt, 0.1 g of TiO₂, 50 g of nickel oxide, and 50 g of ZSM-5 as an inner layer side and 100.1 g of Pt-ZSM and 10 g of SiO₂ as an outer layer side respectively, per liter of carrier.

[0057] Catalyst Element 5 alone was used as a catalyst.

EXAMPLE 5

[0058] A catalyst was prepared according to the procedure of Example 1 (Catalyst Element 1) while changing the amount of nickel oxide to 200 g. The resulting catalyst element (Catalyst Element 6) contained 0.01 g of Pt, 0.1 g of TiO₂, 10 g of nickel oxide, and 50 g of ZSM-5, per liter of carrier.

[0059] Disposing Catalyst Element 6 on the former stage and Catalyst Element 2 on the latter stage formed a catalyst.

EXAMPLE 6

[0060] A catalyst was prepared according to the procedure of Example 1 (Catalyst Element 1) while changing the amount of the ZSM-5 type zeolite to 216 g. The resulting catalyst element (Catalyst Element 7) contained 0.01 g of Pt, 0.1 g of TiO₂, 50 g of nickel oxide, and 10 g of ZSM-5, per liter of carrier.

[0061] Disposing Catalyst Element 7 on the former stage and Catalyst Element 2 on the latter stage formed a catalyst.

EXAMPLE 7

[0062] A catalyst was prepared according to the procedure of Example 1 (Catalyst Element 1) while changing the amount of nickel oxide to 200 g and the amount of the ZSM-5 type zeolite to 216 g, respectively. The resulting catalyst element (Catalyst Element 8) contained 0.01 g of Pt, 0.1 g of TiO₂, 10 g of nickel oxide, and 10 g of ZSM-5, per liter of carrier.

[0063] Disposing Catalyst Element 8 on the former stage and Catalyst Element 2 on the latter stage formed a catalyst.

EXAMPLE 8

[0064] A catalyst was prepared according to the procedure of Example 1 (Catalyst Element 1) while using α -hydrated iron oxide instead of the nickel oxide. The α -hydrated iron oxide was prepared in accordance with the report (Ishikawa et al., Journal of Japan Chemical Society, 91, 935, 1970). Specifically, the α -hydrated iron oxide was obtained by completely dissolving 120 g of iron sulfate (the reagent grade, available from Wako Pure Chemical Industries Ltd. in Japan) in 500 ml of deionized water, gradually adding dropwise an aqueous 1.5 mol/L sodium hydroxide solution (the reagent grade, available from Wako Pure Chemical Industries Ltd. in Japan) to the aqueous iron sulfate solution, adjusting the produced solution to a pH of about 3, changing the temperature of the solution to about 50°C, allowing the solution to remain in a stirred state for about 30 hours, separating the produced precipitate by filtration, thoroughly washing the separated precipitate with water, and drying the washed precipitate at 120°C.

[0065] The resulting catalyst element (Catalyst Element 9) contained 0.01 g of Pt, 0.1 g of TiO₂, 10 g of α -hydrated iron oxide, and 50 g of ZSM-5, per liter of carrier.

[0066] Disposing Catalyst Element 9 on the former stage and Catalyst Element 2 on the latter stage formed a catalyst.

EXAMPLE 9

[0067] A catalyst was prepared according to the procedure of Example 3 (Catalyst Element 4) while using α -hydrated iron oxide instead of the nickel oxide. The resulting catalyst element (Catalyst Element 10) contained 0.01 g of Pt, 0.1 g of TiO₂, 50 g of α -hydrated iron oxide, 50 g of ZSM-5, and 100.1 g of Pt-ZSM, per liter of carrier.

[0068] Catalyst Element 10 alone was used as a catalyst.

EXAMPLE 10

[0069] A catalyst was prepared according to the procedure of Example 4 (Catalyst Element 5) while using α -hydrated iron oxide instead of the nickel oxide. The resulting catalyst element (Catalyst Element 11) contained 0.01 g of Pt, 0.1 g of TiO₂, 50 g of α -hydrated iron oxide, and 50 g of ZSM-5 on the inner side layer and 100.1 g of Pt-ZSM and 10 g of SiO₂ on the outer side layer, respectively per liter of carrier.

[0070] Catalyst Element 11 alone was used as a catalyst.

COMPARATIVE EXAMPLE 1

[0071] Catalyst Element 1 alone was used as a catalyst.

COMPARATIVE EXAMPLE 2

[0072] Catalyst Element 2 alone was used as a catalyst.

COMPARATIVE EXAMPLE 3

[0073] A catalyst was prepared according to the procedure of Example 1 (Catalyst Element 1) while omitting the ZSM-5 type zeolite. The resulting catalyst element (Catalyst Element 12) contained 0.01 g of Pt, 0.1 g of TiO₂, and 50 g of nickel oxide, per liter of carrier.

[0074] Disposing Catalyst Element 12 on the former stage and Catalyst Element 2 on the latter stage formed a catalyst.

COMPARATIVE EXAMPLE 4

[0075] A catalyst was prepared according to the procedure of Example 1 (Catalyst Element 1) while omitting the nickel oxide. The resulting catalyst element (Catalyst Element 13) contained 0.01 g of Pt, 0.1 g of TiO₂, and 50 g of ZSM-5, per liter of carrier.

[0076] Disposing Catalyst Element 13 on the former stage and Catalyst Element 2 on the latter stage formed a catalyst.

COMPARATIVE EXAMPLE 5

[0077] A catalyst was prepared according to the procedure of Example 1 (Catalyst Element 1) while omitting Powder 1. The resulting catalyst element (Catalyst Element 14) contained 50 g of nickel oxide and 50 g of ZSM-5, per liter of carrier.

[0078] Disposing Catalyst Element 14 on the former stage and Catalyst Element 2 on the latter stage formed a catalyst.

EXAMPLE 10

[0079] The catalysts obtained in the examples and the comparative examples described above were tested for the ability to purify the exhaust gas by the following method.

[0080] In the test, a supercharging antechamber combustion type diesel engine (4 cylinders, 2.8 L) was used as the internal combustion engine, and light oil having a sulfur content of 0.05 wt. % used as the fuel for the internal combustion engine.

[0081] First, a given exhaust gas purifying catalyst was set at place in an exhaust gas tube connected to the diesel engine mentioned above and a temperature elevation test from 150°C to 350°C (20°C/min) was carried out by a programmed operation. The programmed operation was performed by the use of an automatic engine-operating device (available from Horiba Seisakusho Ltd. in Japan and sold under the product code of "HERT-381" under the condition of 1500 rpm), with the magnitude of automatic torque varied.

[0082] Then, the programmed operation was performed in the absence of the catalyst from the exhaust gas tube. The gas then exhausted from the engine was continuously sampled and analyzed with a continuous gas analyzer to determine NO_x and hydrocarbon contents in the exhaust gas. NO_x was analyzed with a chemical emission spectral analyzer (CLD) and the hydrocarbon with a hydrogen flame ionizing analyzer (NDIR) respectively. The analysis results at various temperatures are shown in Table 1.

TABLE 1

Gas temperature	150°C	175°C	200°C	250°C	300°C	350°C
Rotational frequency (rpm)	1500	←	←	←	←	←
THC(ppm)	68	60	57	55	35	32
NO _x (ppm)	70	72	82	83	85	95

TABLE 1 (continued)

THC/NO _x	0.97	0.83	0.70	0.66	0.41	0.34
---------------------	------	------	------	------	------	------

[0083] Then, the exhaust gas that had contact with the catalyst was analyzed to determine NO_x and hydrocarbon contents. The purification ratios (the degrees of conversion) of NO_x and hydrocarbon were calculated based on the contents of NO_x and hydrocarbon in the exhaust gas determined in the absence of the catalyst from the exhaust gas tube and the contents of NO_x and hydrocarbon determined in the presence of the catalyst in the exhaust gas tube. Let XO (mol) stand for the content in the exhaust gas in the absence of the catalyst and X1 for the content in the presence of the catalyst, the calculation will be attained by the following formula:

Purification ratio (degree of conversion) (%) =

$$\{(XO - X1)/XO\} \times 100 .$$

[0084] The capacity for purifying NO_x (degree of purification) at various temperatures found by the calculation is shown in Table 2.

TABLE 2

(Purification ratio, %)						
	150°C	175°C	200°C	250°C	300°C	350°C
Ex 1	38	30	40	29	12	10
Ex 2	35	25	35	25	10	9
Ex 3	28	20	20	15	8	8
Ex 4	18	15	20	18	10	9
Ex 5	15	12	30	28	13	5
Ex 6	40	30	25	18	14	8
Ex 7	14	13	25	28	14	8
Ex 8	30	25	25	22	12	11
Ex 9	21	16	15	14	10	10
Ex 10	15	12	14	22	10	11
Com Ex 1	40	32	15	-40	-10	10
Com Ex 2	5	8	20	30	15	5
Com Ex 3	40	28	32	8	6	6
Com Ex 4	3	5	18	28	14	5
Com Ex 5	32	25	32	25	10	10
Ex: Example						
Com Ex: Comparative Example						

[0085] Fig. 1 shows a special property in the adsorption and desorption of NO_x and hydrocarbon when the catalyst formed solely of Catalyst Element 1 obtained in Comparative Example 1 was tested. The data confirm that the catalyst adsorbed the NO_x and the hydrocarbon when the temperature was low, and released the adsorbed NO_x and hydrocarbon simultaneously when the gas temperature rose. The combination of the adsorbent manifesting such special property in the adsorption and desorption of the NO_x and the hydrocarbon with a NO_x purifying catalyst was evidently effective in reducing the NO_x over the range of low, exhaust gas temperature, as compared with the sole use of the NO_x purifying catalyst (Fig. 2).

[0086] Fig. 3 shows the purification ratios of NO_x attained by the catalysts of Example 1 and Comparative Examples 3 and 4. In Comparative Example 3 the hydrocarbon adsorbent was omitted from Example 1, and in Comparative Example 4 the NO_x adsorbent was omitted from Example 1. Comparative Example 3 produced the result of a lower

ratio of NO_x purification than Example 1, because the shortage of the hydrocarbon as a reducing agent in the temperature range permitted the NO_x emission. In contrast, Comparative Example 4 produced a low ratio of NO_x purification in low temperatures because the NO_x adsorbent was omitted. These results clearly indicate that the NO_x purification over a wide range of temperature necessitates the combination of a NO_x adsorbent with a hydrocarbon adsorbent.

[0087] Incidentally, the amounts of catalyst component deposited were expressed based on the apparent volume of the carrier. They may be substantially equal when they are expressed based on the catalyst.

Claims

1. An exhaust gas purifying catalyst, **characterized by** comprising a first catalyst component containing a refractory inorganic oxide carrying a platinum family metal, a nitrogen oxide adsorbent, and a hydrocarbon adsorbent and a second catalyst component for the purification of nitrogen oxide.
2. A catalyst according to claim 1, wherein said first catalyst component said nitrogen oxide adsorbent includes nickel or hydrated iron oxide, said hydrocarbon adsorbent includes zeolite, and said platinum family metal is at least one member selected from the group consisting of platinum, palladium, and rhodium.
3. A catalyst according to claim 1 or claim 2, wherein said first catalyst component is disposed at a high concentration on the upstream side and said second catalyst component is disposed at a high concentration on the downstream side relative to the flow direction of the exhaust gas.
4. A catalyst according to any of claims 1 to 3, wherein said catalyst is a set of at least two pieces and said first catalyst component is disposed on the upstream side and said second catalyst component on the downstream side respectively relative to the flow direction of the exhaust gas.
5. A catalyst according to claim 1 or claim 2, wherein said first and second catalyst components are uniformly mixed.
6. A catalyst according to claim 1 or claim 2, wherein said first catalyst component is made to form an inner layer and said second catalyst component is disposed on the outside of the inner layer.
7. A method for the purification of an exhaust gas, **characterized by** forwarding an exhaust gas having the molar ratio of hydrocarbon to nitrogen oxide (hydrocarbon: nitrogen oxide) in the range of 0.1 to 2 : 1 into contact with an exhaust gas purifying catalyst set forth in any of claims 1 to 6.
8. A method according to claim 7, wherein said exhaust gas is produced from a diesel engine.

FIG. 1

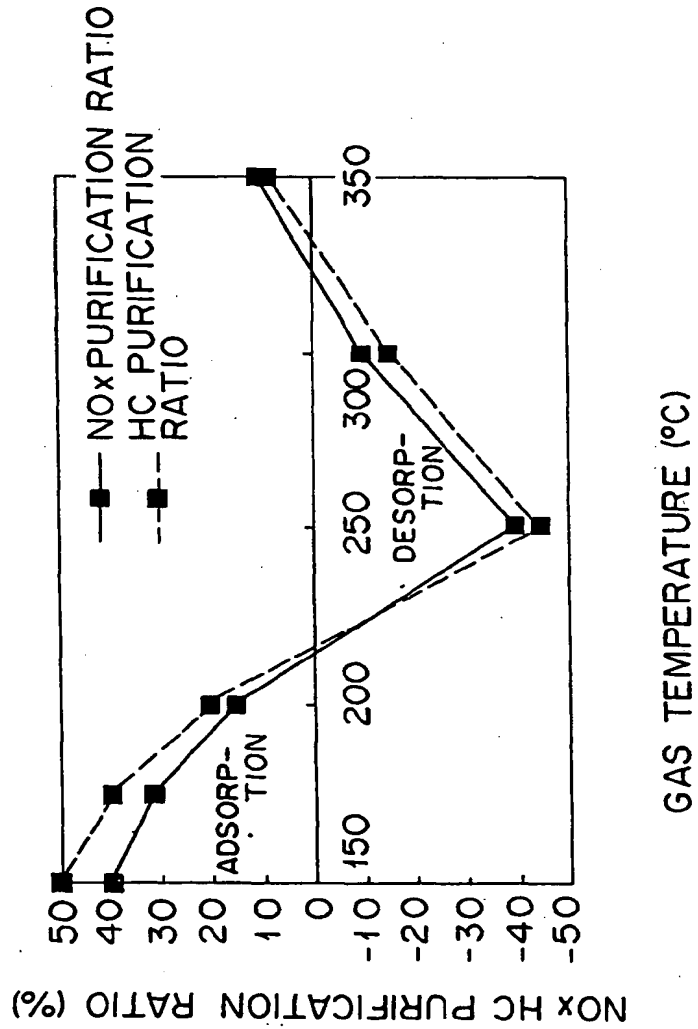


FIG. 2

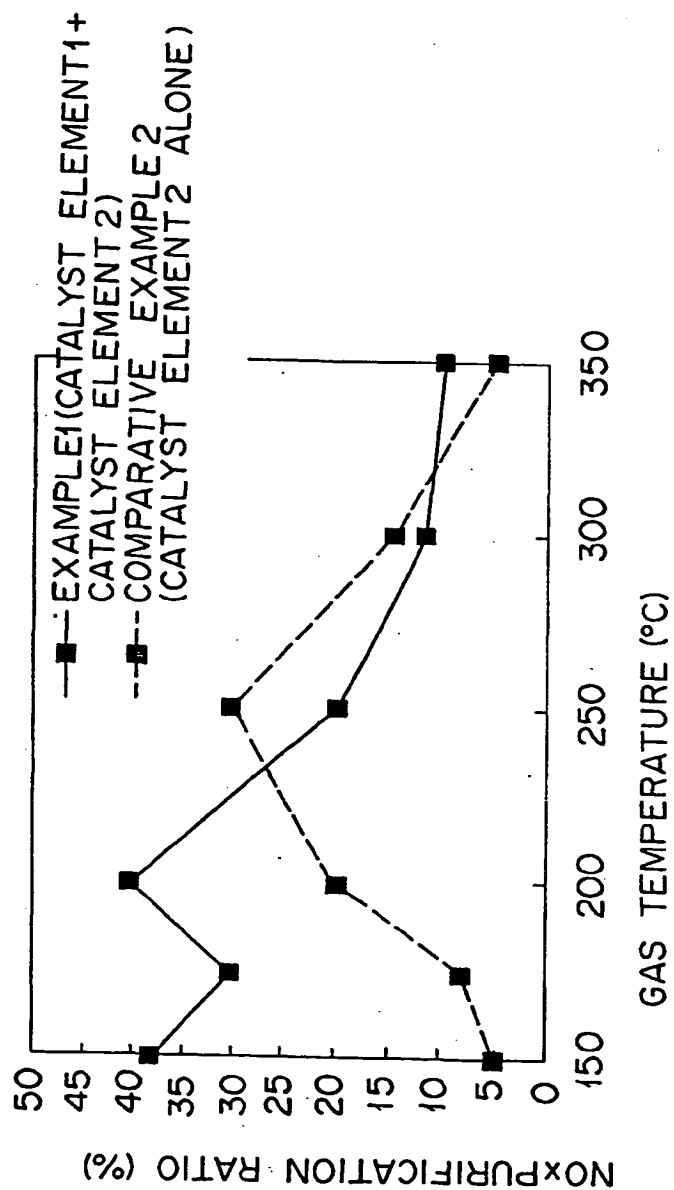
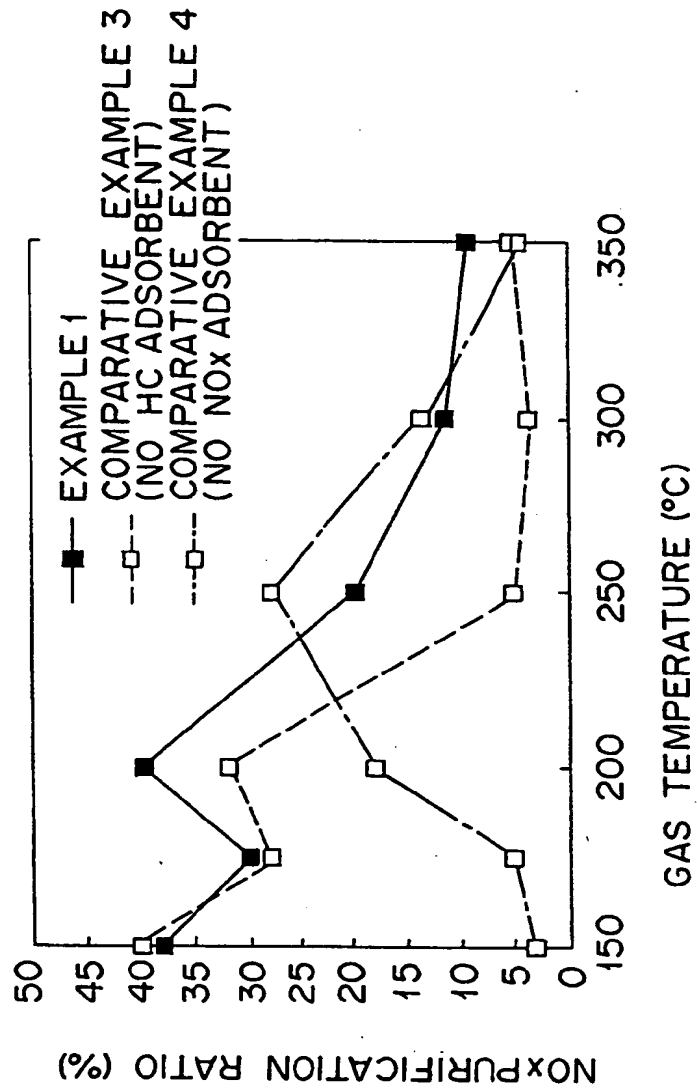


FIG. 3





European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 01 11 4550

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	EP 0 984 142 A (TOYOTA MOTOR CO LTD) 8 March 2000 (2000-03-08) * column 4, line 12 - column 5, line 22; figures 1-4,9 *	1-4,7,8	B01D53/94
X	EP 0 749 774 A (NGK INSULATORS LTD) 27 December 1996 (1996-12-27) * claims; figures 13-33; examples *	1,3,4,7,8	
X	US 5 804 526 A (NARISHIGE TAKESHI ET AL) 8 September 1998 (1998-09-08) * column 3, line 21 - line 26; claims 1-5; figure 1 *	1,6,7	
X	US 5 958 826 A (KUROKAWA TAKAHIRO ET AL) 28 September 1999 (1999-09-28) * abstract; claims 1-8; table 9 *	1,5,7	
X	PATENT ABSTRACTS OF JAPAN vol. 1997, no. 02, 28 February 1997 (1997-02-28) & JP 08 281110 A (NISSAN MOTOR CO LTD), 29 October 1996 (1996-10-29) * abstract *	1-4,6-8	
X	PATENT ABSTRACTS OF JAPAN vol. 1997, no. 02, 28 February 1997 (1997-02-28) & JP 08 281111 A (NISSAN MOTOR CO LTD), 29 October 1996 (1996-10-29) * abstract *	1-4,6-8	TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			B01D F01N
The present search report has been drawn up for all claims			
Place of search MUNICH		Date of completion of the search 27 September 2001	Examiner Eijkenboom, A
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1603 03.82 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 01 11 4550

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

27-09-2001

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
EP 0984142	A	08-03-2000	EP	0984142 A1	08-03-2000
			WO	9848153 A1	29-10-1998
EP 0749774	A	27-12-1996	JP	9000872 A	07-01-1997
			EP	0749774 A2	27-12-1996
			EP	0908225 A2	14-04-1999
			US	5884473 A	23-03-1999
US 5804526	A	08-09-1998	JP	7163871 A	27-06-1995
			CA	2150376 A1	06-04-1995
			DE	69424766 D1	06-07-2000
			DE	69424766 T2	28-09-2000
			EP	0671208 A1	13-09-1995
			WO	9509047 A1	06-04-1995
US 5958826	A	28-09-1999	JP	9010601 A	14-01-1997
			DE	19617123 A1	31-10-1996
			DE	19617124 A1	31-10-1996
			JP	9010594 A	14-01-1997
			US	5795840 A	18-08-1998
			US	5677258 A	14-10-1997
JP 08281110	A	29-10-1996	NONE		
JP 08281111	A	29-10-1996	NONE		

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

THIS PAGE BLANK (USPTO)